

**Amendment to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:**

1. (Currently Amended) A method for diagnosis of a disorder associated with the development of beta amyloid deposits or fibrils in a human or animal subject or assessing the efficacy of treatment rendered to the subject for such disorder, said method comprising the steps step of:
  - A) determining the presence of mtDNA CR mutations; and
  - B) comparing a mtDNA CR value obtained by the quantitative determination made in Step A with a mtDNA CR value representative of subjects who suffer from a disorder associated with the development of beta amyloid deposits or fibrils.
2. (Currently Amended) A method according to Claim 1, wherein Step A comprises making a qualitative determination that mtDNA mtDNS CR mutation is or is not present.
3. (Currently Amended) A method according to Claim 1, wherein Step A comprises making a quantitative determination of mtDNA CR mtDNS mutations.
4. (Currently Amended) A method according to Claim 3 further comprising the step of:
  - B) comparing a mtDNA mtDNS CR value obtained by the quantitative determination made in Step A with a control mtDNA mtDNS CR value to determine whether the subject has significantly more mtDNA mtDNS CR mutations than control.
5. (Cancelled)
6. (Original) A method according to any of Claim 1 wherein Step A comprises testing for a

T4141G mutation.

7. (Withdrawn) A method according to any of Claim 1 wherein Step A comprises testing for a T414C mutation.
8. (Withdrawn) A method according to any of Claim 1 wherein Step A comprises testing for a T477C mutation.
9. (Withdrawn) A method according to any of Claim 1 wherein Step A comprises testing for a T146C mutation.
10. (Withdrawn) A method according to any of Claim 1 wherein Step A comprises testing for a T152C mutation.
11. (Withdrawn) A method according to any of Claim 1 wherein Step A comprises testing for a A189G mutation.
12. (Withdrawn) A method according to any of Claim 1 wherein Step A comprises testing for a T195C mutation.
13. (Original) A method according to Claim 1 wherein Step A is carried out at least in part by PNA-clamping PCR.
14. (Original) A method according to Claim 1 wherein Step A is carried out at least in part by oligonucleotide hybridization.
15. (Original) A method according to Claim 1 wherein Step A is carried out at least in part by primer extension.

16. (Original) A method according to Claim 1 wherein Step A is carried out at least in part by restriction digestion.
17. (Original) A method according to Claim 1 wherein the determination of Step A is made in a specimen of tissue, cells or body fluid selected from the group consisting of:
  - i. brain tissue;
  - ii. brain tissue from the frontal cortex;
  - iii. nervous tissue;
  - iv. nerve cells
  - v. blood
  - vi. blood cells;
  - vii. urine;
  - viii. urinary tract cells;
  - ix. skin;
  - x. skin cells;
  - xi. epithelium;
  - xii. epithelial cells;
  - xiii. fibroblasts;
  - xiv. cerebrospinal fluid; and
  - xv. cells contained in cerebrospinal fluid.
18. (Original) A method according to Claim 1 wherein the method is carried out for post-symptomatic diagnosis of a disorder in a subject who has begun to exhibit symptoms of that disorder.
19. (Original) A method according to Claim 1 wherein the method is carried out for pre-symptomatic diagnosis of a disorder in a subject who has not begun to exhibit symptoms of that disorder.
20. (Original) A method according to Claim 1 wherein the disorder is a neurodegenerative disease.

21. (Original) A method according to Claim 1 wherein the disorder is Alzheimer's Disease.
22. (Withdrawn) A method according to Claim 1 wherein the disorder is Parkinson's Disease.
23. (Withdrawn) A method according to Claim 1 wherein the disorder is Down's Syndrome-associated dementia.
24. (Withdrawn) A method according to Claim 1 wherein the disorder is a spongiform encephalopathy.
25. (Withdrawn) A method according to Claim 1 wherein the disorder is type II diabetes .
26. (Withdrawn) A method according to Claim 1 wherein the disorder is Creutzfeldt-Jakob disease.
27. (Withdrawn) A method according to Claim 1 wherein the disorder is a Huntington's disease.
28. (Withdrawn) A method according to Claim 1 wherein the disorder is macular degeneration.
29. (Withdrawn) A method according to Claim 1 wherein the disorder is a prion disease.
30. (Withdrawn) A method according to Claim 1 wherein Step A comprises:
  - obtaining sample cells from the subject;
  - extracting DNA from the sample cells;
  - subjecting the extracted DNA to mitochondrial DNA control region amplification;
  - determining whether homoplasmic 414 and 477 nucleotide variants are present by direct sequencing for heteroplasmic 414 and 477 nucleotide mutations; and

if 414 and 477 nucleotide variants are detected, cloning the mutant molecules and sequencing the clone.

31. (Cancelled)

32. (Cancelled)

33. (Cancelled)

34. (Cancelled)